### **CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

### **Listing of Claims:**

Claim 1 (currently amended): A compound represented by formula I:

$$R^2$$
 $R^3$ 
 $Q$ 
 $Q$ 
 $R^4$ 
 $Q$ 
 $Q$ 

wherein

 $\mathbf{R}^1$  is selected from the group consisting of H, halogen,  $(C_{1-4})$ alkyl,  $O(C_{1-6})$ alkyl, and haloalkyl;

 $R_2$  is H or  $(C_{1-4})$ alkyl;

 $\mathbb{R}^3$  is H or  $(C_{1-4})$ alkyl;

 $\mathbf{R}^4$  is  $(C_{1-4})$ alkyl,  $(C_{1-4})$ alkyl $(C_{3-7})$ cycloalkyl, or  $(C_{3-7})$ cycloalkyl; and

**Q** is a fused phenyl-5 or 6-membered saturated heterocycle having one to two heteroatoms selected from O and N, said **Q** is selected from the group consisting of:

R<sup>7</sup> R<sup>7</sup>

a) wherein one of **E** and **G** is C(O) and the other is  $NR^5$  wherein  $R^5$  is selected from the group consisting of H, hydroxy and  $(C_{1-4})$ alkyl unsubstituted or substituted with pyridinylmethyl, (pyridinyl-N-oxide)methyl or  $C(O)OR^6$  wherein  $R^6$ 

is H or  $(C_{1-4})$ alkyl; and each  $\mathbb{R}^7$  is independently H, Me or Et; or

b) wherein **E** is NR<sup>8</sup> wherein R<sup>8</sup> is H, (C<sub>1-4</sub>)alkyl unsubstituted or substituted with C(O)OR<sup>9</sup> wherein R<sup>9</sup> is H or (C<sub>1-4</sub>)alkyl; or

c) wherein **D** and **G** are  $NR^{10}$  wherein each  $R^{10}$  is independently H or  $(C_{1-4})$ alkyl unsubstituted or substituted with  $C(O)OR^{11}$  wherein  $R^{11}$  is H or  $(C_{1-4})$ alkyl; or

d) wherein one of **L**, **M**, **Y** and **Z** is  $NR^{12}$  wherein  $R^{12}$  is H,  $(C_{1-4})$ alkyl unsubstituted or substituted with  $C(O)OR^{12x}$  wherein  $R^{12x}$  is H or  $(C_{1-4})$ alkyl; one of the remaining positions of **L**, **M**, **Y** and **Z** adjoining the  $NR^{12}$  is C(O); and the remaining two positions are each  $CR^{13}R^{13}$  wherein each  $R^{13}$  is independently H, Me or Et; or

e) wherein three adjoining positions of **L**, **M**, **Y** and **Z** (namely **L-M-Y** or **M-Y-Z**) represent  $NR^{14}$ -C(O)-O- or  $-NR^{15}$ -C(O)- $NR^{16}$  — wherein  $R^{14}$ ,  $R^{15}$  and  $R^{16}$  each represents H or  $(C_{1-4})$ alkyl unsubstituted or substituted with C(O)O $R^{17}$  wherein  $R^{17}$  is H or  $(C_{1-4})$ alkyl; and the remaining position of **L**, or **Z** is  $CR^{18}R^{18}$  wherein each  $R^{18}$  is H, Me or Et;

or a pharmaceutically acceptable salt, or prodrug thereof.

Claim 2 (currently amended): The compound according to claim 1, wherein  $\mathbf{R}^1$  is

selected from: H, Cl, F,  $(C_{1-4})$ alkyl and  $CF_3$ ;  $\mathbf{R}^2$  and  $\mathbf{R}^3$  is each independently H or Me;  $\mathbf{R}^4$  is ethyl or cyclopropyl; and

# **Q** is selected from:

$$N-R^5$$
 or  $R^5$  wherein  $R^5$  is H, hydroxy,  $CH_3$  or (4-

## pyridinyl)methyl;

$$N-Me$$
 $N-CMe_3$ 
 $Me$ 
 $N-CMe$ 
 $N-CMe$ 

or CH<sub>2</sub>C(O)OH,

# or **Q** is further selected from:

$$R^{18}$$
  $R^{18}$   $R$ 

CH<sub>2</sub>C(O)OH and each  $\mathbb{R}^{18}$  is independently H or Me<sub>2</sub>. More preferably,  $\mathbb{R}^{14}$  is H or CH<sub>2</sub>C(O)OH and each  $\mathbb{R}^{18}$  is H,

or Q is further selected from:

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wherein R<sup>15</sup> is H, Me or CH<sub>2</sub>C(O)OH and

 $\mathbf{R}^{16}$  is H, Me or CH<sub>2</sub>C(O)OH. More preferably,  $\mathbf{R}^{15}$  is H or CH<sub>2</sub> and  $\mathbf{R}^{16}$  is H, CH<sub>3</sub> or CH<sub>2</sub>-C(O)OH.

Claim 3 (original): The compound according to claim 2, wherein  $\mathbb{R}^1$  is H, Cl, F or Me;  $\mathbb{R}^2$  is H;  $\mathbb{R}^3$  is Me;  $\mathbb{R}^4$  is ethyl; and  $\mathbb{Q}$  is selected from:

$$N-R^5$$

wherein **R**<sup>5</sup> is H, hydroxy or (4-pyridinyl)methyl;

N-Me N-Me or 
$$R^{18}$$
  $R^{18}$   $R^{18}$ 

wherein  $\mathbf{R}^{14}$  is H or

 $CH_2C(O)OH$  and each  $\mathbf{R}^{18}$  is H,

or

$$R^{15}$$
 or  $R^{15}$   $R^{16}$   $R^{16}$ 

wherein  $\mathbf{R}^{15}$  is H or CH<sub>3</sub> and  $\mathbf{R}^{16}$  is H, CH<sub>3</sub>

or CH<sub>2</sub> C(O)OH.

Claim 4 (original): The compound according to claim 3, wherein **Q** is selected from:

Claim 5 (original): The compound according to claim 4, wherein  $\mathbb{R}^1$  is H,  $\mathbb{R}^2$  is H,  $\mathbb{R}^3$  is Me,  $\mathbb{R}^4$  is ethyl and  $\mathbb{Q}$  is selected from:

Claim 6 (currently amended): A pharmaceutical composition for the treatment or prevention of HIV infection, comprising a compound of formula I according to claim 1, or a pharmaceutically acceptable salt, or prodrug thereof, and a pharmaceutically acceptable carrier.

Claim 7 (currently amended): A method for the treatment or prevention of HIV infection, comprising administering to a patient an HIV inhibiting amount of a compound of formula I according to claim 1, or a pharmaceutically acceptable salt, or

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prodrug thereof.

Claim 8 (currently amended): A method for the treatment or prevention of HIV infection, comprising administering to a patient an HIV inhibiting amount of a pharmaceutical composition, according to claim 6.

Claim 9 (cancelled)

Claim 10 (currently amended): A method for preventing perinatal transmission of HIV11-HIV-1 from mother to baby, comprising administering a compound of formula I according to claim 1, to the mother before giving birth.

Claim 11 (cancelled)

Claim 12 (original): A process for producing a compound of formula I according to claim 1, comprising steps of:

- coupling a compound of formula 2:

wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are as defined in claim 1;

with a phenolic derivative selected from:

HO 
$$\stackrel{\mathsf{R}^7}{=}_{\mathsf{G}}^{\mathsf{R}^7}$$
 wherein one of **E** and **G** is C(O) and the other is N**R**<sup>5A</sup>

wherein one of **E** and **G** is C(O) and the other is N**R**<sup>5A</sup> wherein  $\mathbf{R}^{5A}$  is a N-protecting group, hydroxy or (C<sub>1-4</sub>)alkyl unsubstituted or substituted with pyridylmethyl, (pyridinyl-N-oxide) methyl or C(O)O**R**<sup>6A</sup> wherein  $\mathbf{R}^{6A}$  is a carboxy protecting group or (C<sub>1-4</sub>)alkyl; and each  $\mathbf{R}^7$  is independently H, Me or Et.

b) wherein **E** is NR<sup>8A</sup> wherein R<sup>8A</sup> is a N-protecting group,  $(C_{1-4})$ alkyl unsubstituted or substituted with  $C(O)OR^{9A}$  wherein R<sup>9A</sup> is a carboxy protecting group or  $(C_{1-4})$ alkyl; or

$$HO \longrightarrow G$$

c) wherein **D** and **G** each independently is  $NR^{10A}$  wherein  $R^{10A}$  is a N-protecting group or  $(C_{1-4})$ alkyl unsubstituted or substituted with  $C(O)OR^{11A}$  wherein  $R^{11A}$  is a carboxy protecting group or  $(C_{1-4})$ alkyl;

d) wherein one of **L**, **M**, **Y** and **Z** is  $NR^{12A}$  wherein  $NR^{12A}$  is a N-protecting group,  $(C_{1-4})$ alkyl unsubstituted or substituted with  $C(O)OR^{12y}$  wherein  $R^{12y}$  is a carboxy protecting group or  $(C_{1-4})$ alkyl; one of the remaining positions of **L**, **M**, **Y** and **Z** adjoining the  $NR^{12A}$  is C(O); and the remaining two positions are each  $CR^{13}R^{13}$  wherein each  $R^{13}$  is independently H, Me or Et; or

e) wherein three adjoining positions of **L**, **M**, **Y** and **Z** (namely **L-M-Y** or **M-Y-Z**) represent -NR<sup>14</sup>-C(O)-O- or -NR<sup>15</sup>-C(O)- NR<sup>16</sup>- wherein R<sup>14</sup>, R<sup>15</sup> and R<sup>16</sup> are as defined in claim 1, and the remaining position of **L** or **Z** is CR<sup>18</sup>R<sup>18</sup> wherein each R<sup>18</sup> is as defined in claim 1; and, if required,

- removing any protective groups in a mixture of aqueous base or aqueous acid in a co-solvent, to obtain the corresponding compound of formula I.

#### Claim 13 (cancelled)

Claim 14 (cancelled)

Claim 15 (cancelled)